

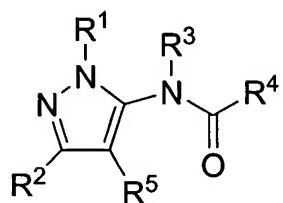
AMENDMENTS TO THE CLAIMS

Please cancel Claims 1-20 and insert therefor Claims 21-35 as follow. This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1-20. (Canceled)

21. (New) A compound of the formula I:



I

wherein:

R¹ is selected from the group consisting of:

- (1) hydrogen,
- (2) C₁-6alkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl,
- (3) C₃-7cycloalkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl, and
- (4) phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:
 - (a) -C₁-6alkyl,
 - (b) -O-C₁-6alkyl,
 - (c) halo,
 - (d) hydroxy,
 - (e) trifluoromethyl,
 - (f) -OCF₃,
 - (g) -CO₂R⁹,

wherein R⁹ is independently selected from:

- (i) hydrogen,
- (ii) -C₁-6alkyl, which is unsubstituted or substituted with 1-6 fluoro,
- (iii) benzyl, and

(iv) phenyl,
(h) -NR¹⁰R¹¹,
wherein R¹⁰ and R¹¹ are independently selected from:
(i) hydrogen,
(ii) -C₁₋₆alkyl, which is unsubstituted or substituted with 1-6
fluoro,
(iii) -C₅₋₆cycloalkyl,
(iv) benzyl,
(v) phenyl,
(vi) -S(O)₂-C₁₋₆alkyl,
(vii) -S(O)₂-benzyl, and
(viii) -S(O)₂-phenyl,
(i) -CONR¹⁰R¹¹, and
(j) -NO₂;

(5) heterocycle, wherein heterocycle is selected from:
benzimidazolyl, benzimidazolonyl, benzofuranyl, benzofurazanyl,
benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl,
carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyl,
indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl,
naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl,
pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl,
pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrahydropyranyl,
tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidinyl,
1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyridin-2-onyl,
pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzimidazolyl,
dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl,
dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl,
dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl,
dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl,
dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl,
dihydrothienyl, dihydrotriazolyl, dihydroazetidinyl, methylenedioxybenzoyl,
tetrahydrofuranyl, and tetrahydrothienyl, and N-oxides thereof, which is
unsubstituted or substituted with one or more substituents independently
selected from:
(a) -C₁₋₆alkyl,
(b) -O-C₁₋₆alkyl,
(c) halo,

- (d) hydroxy,
- (e) phenyl,
- (f) trifluoromethyl,
- (g) -OCF₃,
- (h) -CO₂R⁹,
- (i) -NR¹⁰R¹¹, and
- (j) -CONR¹⁰R¹¹;

R² is phenyl;

R³ is independently selected from the group consisting of:

- (1) hydrogen, and
- (2) C₁-6alkyl;

R⁴ is selected from the group consisting of:

- (1) C₁-6alkyl, which is unsubstituted or substituted with halogen, hydroxyl, phenyl or heterocycle,
- (2) C₃-7cycloalkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl, and
- (3) phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:
 - (a) -C₁-6alkyl,
 - (b) -O-C₁-6alkyl,
 - (c) halo,
 - (d) hydroxy,
 - (e) trifluoromethyl,
 - (f) -OCF₃,
 - (g) -CO₂R⁹,
 - (h) -CN,
 - (i) -NR¹⁰R¹¹,
 - (j) -CONR¹⁰R¹¹, and
 - (k) -NO₂;
- (4) heterocycle, wherein heterocycle is selected from:
benzimidazolyl, benzimidazolonyl, benzofuranyl, benzofurazanyl,
benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl,
carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyl,
indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl,
naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl,

pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrahydropyranyl, tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidinyl, 1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyridin-2-onyl, pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzoimidazolyl, dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl, dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl, dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl, dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl, dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl, dihydrothienyl, dihydrotriazolyl, dihydroazetidinyl, methylenedioxybenzoyl, tetrahydrofuranyl, and tetrahydrothienyl, and N-oxides thereof, which is unsubstituted or substituted with one or more substituents independently selected from:

- (a) -C₁₋₆alkyl,
- (b) -O-C₁₋₆alkyl,
- (c) halo,
- (d) hydroxy,
- (e) phenyl,
- (f) trifluoromethyl,
- (g) -OCF₃,
- (h) -CO₂R⁹,
- (i) -NR¹⁰R¹¹, and
- (j) -CONR¹⁰R¹¹;

R⁵ is independently selected from the group consisting of:

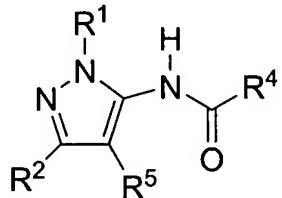
- (1) hydrogen,
- (2) C₁₋₆alkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl,
- (3) C₃₋₇cycloalkyl, which is unsubstituted or substituted with halogen, hydroxyl or phenyl, and
- (4) phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:
 - (a) -C₁₋₆alkyl, which is unsubstituted or substituted with -NR¹⁰R¹¹,
 - (b) -O-C₁₋₆alkyl,
 - (c) halo,

- (d) hydroxy,
- (e) trifluoromethyl,
- (f) $-\text{OCF}_3$;
- (g) $-\text{CO}_2\text{R}^9$,
- (h) $-\text{NR}^{10}\text{R}^{11}$,
- (i) $-\text{C}(\text{O})\text{NR}^{10}\text{R}^{11}$, and
- (j) $-\text{NO}_2$,

(5) heterocycle, wherein heterocycle is selected from:
benzimidazolyl, benzimidazolonyl, benzofuranyl, benzofurazanyl,
benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl,
carbolinyl, cinnolinyl, furanyl, imidazolyl, indolinyl, indolyl, indolazinyl,
indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl,
naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl,
pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl,
pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrahydropyranyl,
tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidinyl,
1,4-dioxanyl, hexahydroazepinyl, piperazinyl, piperidinyl, pyridin-2-onyl,
pyrrolidinyl, morpholinyl, thiomorpholinyl, dihydrobenzimidazolyl,
dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl,
dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl,
dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl,
dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl,
dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl,
dihydrothienyl, dihydrotriazolyl, dihydroazetidinyl, methylenedioxybenzoyl,
tetrahydrofuranyl, and tetrahydrothienyl, and N-oxides thereof, which is
unsubstituted or substituted with one or more substituents independently
selected from:
(a) $-\text{C}_1\text{-}6\text{alkyl}$,
(b) $-\text{O-C}_1\text{-}6\text{alkyl}$,
(c) halo,
(d) hydroxy,
(e) phenyl,
(f) trifluoromethyl,
(g) $-\text{OCF}_3$;
(h) $-\text{CO}_2\text{R}^9$,- (i) $-\text{NR}^{10}\text{R}^{11}$, and
- (j) $-\text{CONR}^{10}\text{R}^{11}$;

or a pharmaceutically acceptable salt thereof.

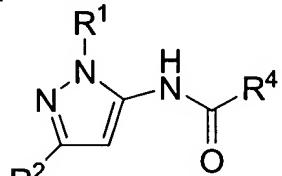
22. (New) The compound of Claim 21 of the formula Ia:



Ia

or a pharmaceutically acceptable salt thereof.

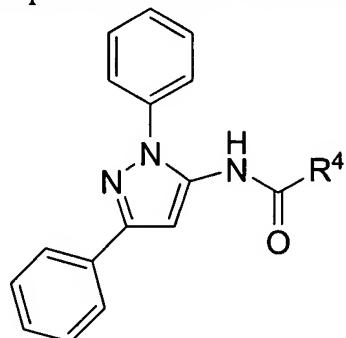
23. (New) The compound of Claim 22 of the formula Ib:



Ib

or a pharmaceutically acceptable salt thereof.

24. (New) The compound of Claim 23 of the formula Ic:



Ic

or a pharmaceutically acceptable salts thereof.

25. (New) The compound of Claim 21 wherein R¹ is phenyl.

26. (New) The compound of Claim 21 wherein R³ is hydrogen.

27. (New) The compound of Claim 21 wherein R⁴ is phenyl, which is unsubstituted or substituted with one or more substituents independently selected from:

- (a) -C₁₋₆alkyl,
- (b) -O-C₁₋₆alkyl,
- (c) halo,
- (d) hydroxy,
- (e) trifluoromethyl,
- (f) -OCF₃;
- (g) -CO₂-C₁₋₆alkyl,
- (h) -CN,
- (i) -NH₂,
- (j) -NH-C₁₋₆alkyl,
- (k) -CONH₂, and
- (l) -CONH-C₁₋₆alkyl.

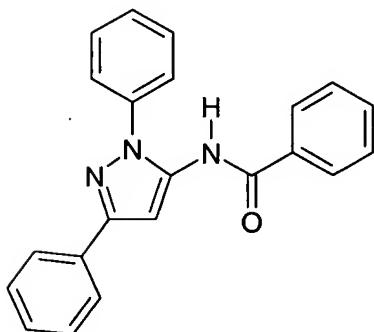
28. (New) The compound of Claim 27 wherein R⁴ is phenyl, which is unsubstituted or substituted with halo or -CN.

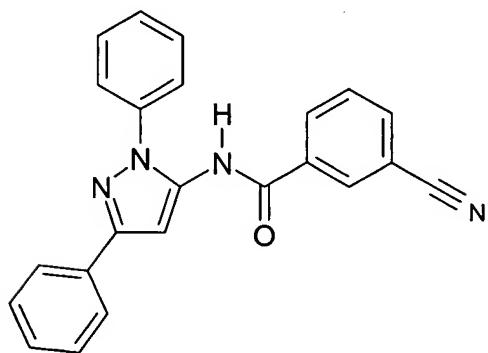
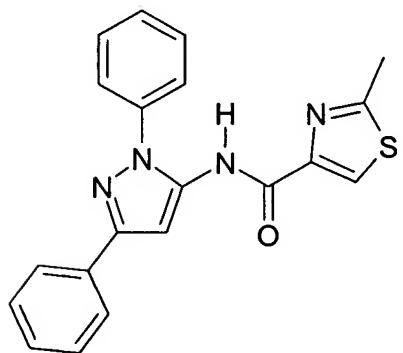
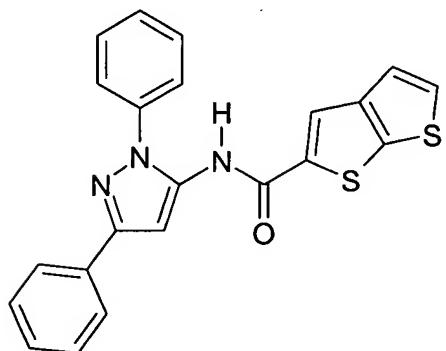
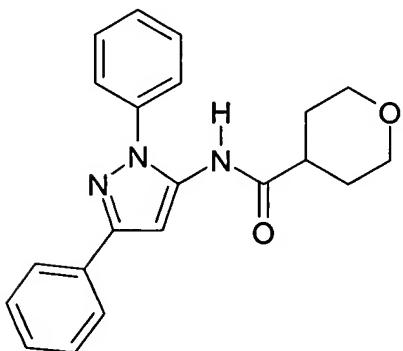
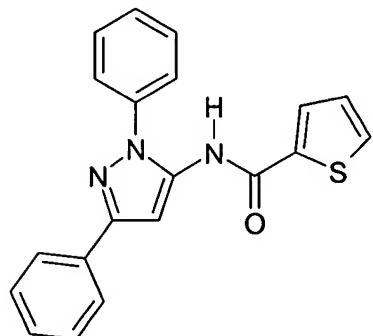
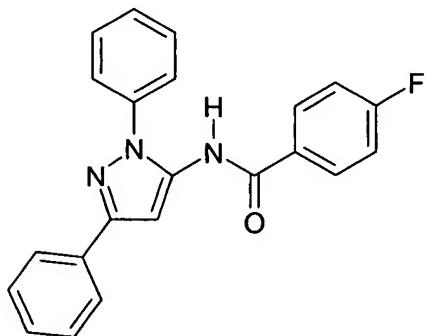
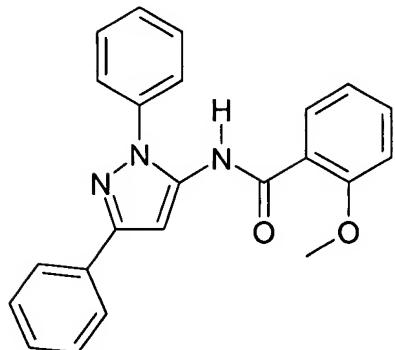
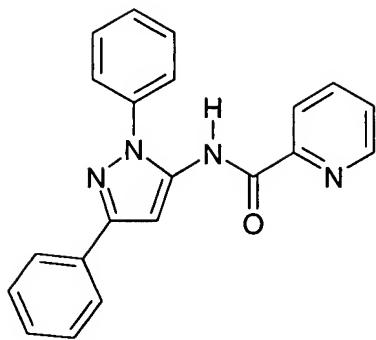
29. (New) The compound of Claim 28 wherein R⁴ is phenyl.

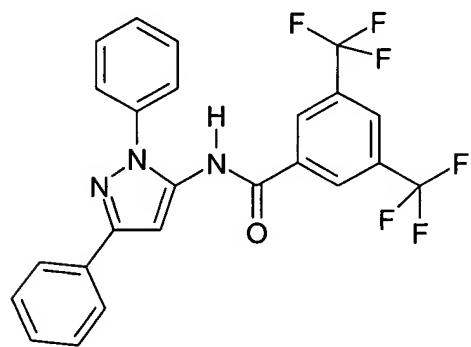
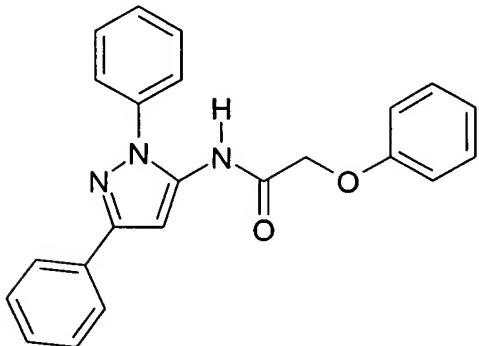
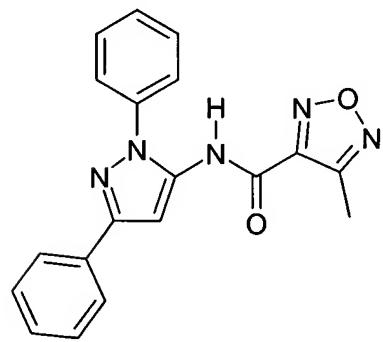
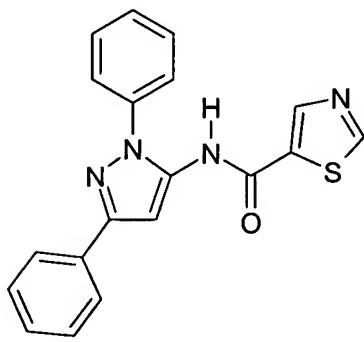
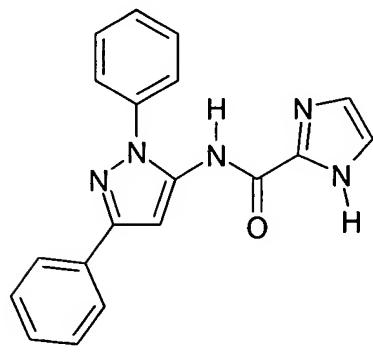
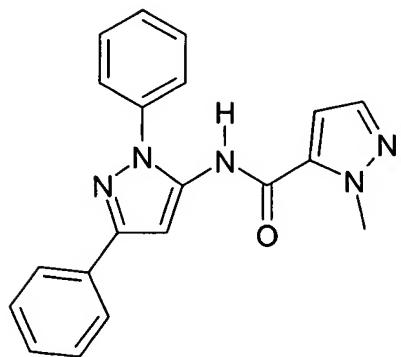
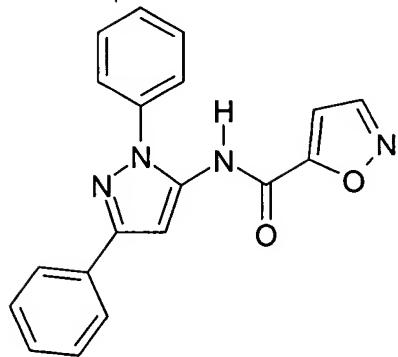
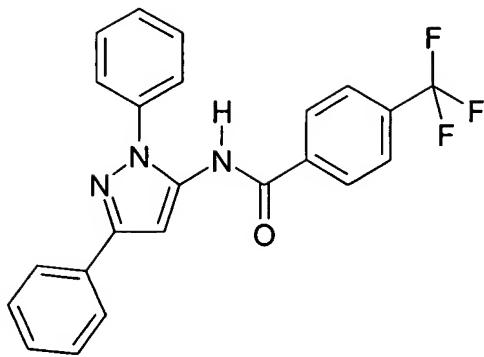
30. (New) The compound of Claim 21 wherein R⁴ is pyridyl.

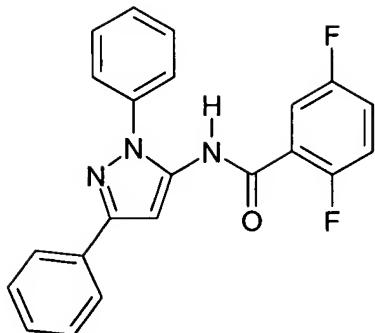
31. (New) The compound of Claim 21 wherein R⁵ is hydrogen.

32. (New) A compound which is selected from the group consisting of:









or a pharmaceutically acceptable salt thereof.

33. (New) A pharmaceutical composition which comprises an inert carrier and the compound of Claim 21 or a pharmaceutically acceptable salt thereof.

34. (New) A method for treating schizophrenia in a mammalian patient in need of such which comprises administering to the patient a therapeutically effective amount of the compound of Claim 21 or a pharmaceutically acceptable salt thereof.

35. (New) A method for treating anxiety in a mammalian patient in need of such which comprises administering to the patient a therapeutically effective amount of the compound of Claim 21 or a pharmaceutically acceptable salt thereof.